

Study of the stability of dried tomato halves during shelf-life to minimise oxidative damage

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Abstract: Optimal operating conditions for storage of dried tomato halves were investigated to obtain decreased oxidative damage, evaluated in terms of colour variation, combined with microbial stability of the product, ie inhibition of growth of eumycetes. Experiments were planned using a saturated factorial design. Variables studied were the moisture content of dried tomato halves in the range 10–60%, temperature in the range 5–30°C and storage time in the range 1–38 days. Eighteen storage experiments were carried out in the dark under vacuum at the storage conditions indicated by the experimental design. At the end of each experiment, surface colour was measured by a tristimulus colorimeter, and growth of eumycetes was evidenced by olfactory and visual perception, followed by qualitative microbial analysis. Eumycetes were present in all stored products, except that at 10% moisture, and, generally speaking, storage conditions did not allow micro-organisms to grow. From isoresponse curves the optimal region for storage to minimise oxidative damage was extrapolated, which was represented by residual moisture values between 20 and 40% and $\leq 18^\circ\text{C}$ storage temperature, with a minimum point of colour variation at approximately 30% moisture content and 10°C .

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Keywords: tomato; oxidative damage; shelf-life; drying

INTRODUCTION

Drying of tomato halves causes oxidative heat damage to the product, shown by both a loss of ascorbic acid and an increase in 5-hydroxymethyl-2-furfural (HMF) content.¹ As a result, undesirable colour and appearance changes occur in dried tomatoes. Conversely, no significant changes in lycopene content occur during drying.¹ This is probably due to both the heat resistance of lycopene^{2–5} and the antioxidant activity of Maillard reaction derivatives.^{6,7}

Optimisation of tomato drying in terms of both maximising the drying rate and minimising oxidative heat damage requires low temperatures for short times (LTST). LTST treatments may be applied by drying tomatoes in small pieces, eg slices, quarters or cubes. A reduction in thickness corresponds to an increase in moisture diffusion, resulting in fast water removal. LTST treatments may also be applied by drying tomato halves, resulting in partial water removal; in this way, intermediate-moisture tomato products can be obtained. In recent years the commercial importance of dried tomato halves has been increasing, since they can be used as a component for several foods such as pizza.

Several studies^{1,5,8} have shown that oxidative damage to dried tomatoes also occurs during storage. In this case, browning has been associated with a

marked lycopene loss of dried tomatoes. There are few data on optimal operating conditions of storage.

The aim of this work was (i) to study the stability of tomato halves during storage as a function of moisture content and time–temperature conditions and (ii) to find the optimal range of storage operating conditions, particularly to minimise oxidative damage.

EXPERIMENTAL

Optimisation procedure

This study was carried out using multivariate statistical analysis.⁹ A saturated factorial design by the central composite design method was first used to evaluate the effect of simultaneous variations in variables. Experiments were then carried out. Finally, a response surface model (RSM) was determined to find both the optimal conditions for the phenomenon studied and the level of individual variables required to reach a particular response level.

Factorial design

Variables studied were the moisture content of dried tomato halves and storage time and temperature.

Moisture content was related to water activity by the following Halsey model, which has already been used

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(Received 10 February 2000; revised version received 28 July 2000; accepted 21 August 2000)

Table 1. Limiting values for variables used in the factorial design

Variable	Minimum value (-)	Maximum value (+)
Moisture content (%)	20	50
Temperature (°C)	10	25
Time (days)	8	30

in other studies on tomatoes:¹⁰

$$a_w = \exp\left(-\frac{B}{n_s^A}\right) \quad (1)$$

where n_s is in gg^{-1} dry matter, $A = 0.93$ and $B = 0.103$. Values for model parameters were determined in a previous study¹ by fitting experimental data of the desorption isotherm of tomato halves.

Since the aim of this work was to find the optimal storage conditions to minimise oxidative damage, variables such as light intensity and oxygen content were not considered owing to their known, relevant effects on oxidative damage to dried tomatoes during storage.^{1,5,8} Experiments were carried out in the dark under vacuum storage conditions.

To apply the saturated factorial design, experimental data were collected using two levels for each of the variables studied. Thus information was obtained which was necessary to determine how the response was influenced by single factors and by their interactions. Table 1 shows the limiting values for the variables used.

The number of experiments necessary for a saturated factorial design by the central composite design method was $2^n + 2n + 4$, where n is the number of variables.⁹ The RSM model requires:

- experiments carried out at the point where all the variables take on their average values (centre point)—the centre point is replicated several times to provide an independent estimate of the experimental error;
- experiments carried out at a distance of 1.68 from the centre point (by adding a pair of experiments along the co-ordinate axes at a distance equal to half the diagonal of the cube);
- experiments carried out at the edges of a cube with the centre point in the middle and with its sides corresponding to each variable range (-1 to $+1$).

Table 2 shows the variable values used for the 18 shelf-life experiments.

Shelf-life experiments

Drying

Ripe, fresh tomatoes of Cencara cultivar (Peviani srl, Milan, Italy) were selected in order to have homogeneous samples in terms of drying time and surface colour.¹

Six drying tests were carried out to obtain samples for the 18 shelf-life experiments.

Thirty tomatoes 5.5 cm in diameter were selected for each drying test. Tomatoes were cut into halves with a knife, and parenchyma and seeds were removed. For each drying test, 40 tomato halves were used, and 20 tomato halves were used to characterise fresh tomatoes in terms of moisture content and surface colour.

Tomato halves were dried in a pilot-plant cabinet air dryer, designed and built by Thermo Lab (Milan, Italy), following the procedure of Zanoni *et al.*¹ The 40 halves were placed on a perforated stainless steel tray (40 cm × 60 cm) in eight rows of five halves. The tray was connected to a balance to measure the tomato weight during drying. Drying was carried out at 80 °C and an air flow rate of 1.5 ms^{-1} by through-flow. Tomatoes were dried to the different final moisture contents reported in Table 2.

At the end of drying, tomato halves were immediately packed under vacuum in several airtight, water-vapour-proof, plastic bags. Bags were immediately chilled at 3 °C for 15 min and then transferred to experimental storage conditions.

For each drying test, 10 dried tomato halves were used to characterise dried tomatoes in terms of surface colour and growth of eumycetes.

Storage

Dried halves were stored in the dark under the time-temperature conditions reported in Table 2, using five thermostatic cells. Samples taken from the cells were characterised in terms of moisture content, surface colour and growth of eumycetes.

Analyses

Moisture content (%) was determined by gravimetry in triplicate following the method described by Zanoni *et al.*¹

Table 2. Operating conditions of shelf-life experiments planned by the factorial design

Experiment	Moisture content (%)	Temperature (°C)	Time (days)
1	10	17.5	19.0
2	60	17.5	19.0
3	35	5.0	19.0
4	35	30.0	19.0
5	35	17.5	0.5
6	35	17.5	37.5
7	20	10.0	8.0
8	50	10.0	8.0
9	20	25.0	8.0
10	50	25.0	8.0
11	20	10.0	30.0
12	50	10.0	30.0
13	20	25.0	30.0
14	50	25.0	30.0
15	35	17.5	19.0
16	35	17.5	19.0
17	35	17.5	19.0
18	35	17.5	19.0

Surface colour of both fresh and dried tomatoes was measured by the Hunter colorimetric system using a Chroma Meter CR-210 tristimulus colorimeter (Minolta Co, Tokyo, Japan). L^* , a^* and b^* parameters of samples were measured, and ΔE values were calculated for dried samples as follows:¹¹

$$\Delta E = \sqrt{(L_t^* - L_0^*)^2 + (a_t^* - a_0^*)^2 + (b_t^* - b_0^*)^2} \quad (2)$$

where L_0^* , a_0^* and b_0^* are the colour parameters of dried tomato halves at time zero of storage and L_t^* , a_t^* and b_t^* are the colour parameters at time t of storage. The colour variation was used as a fast, simple index of oxidative damage.^{12,13} Seven measurements of colour parameters were carried out for each sample.

Growth of eumycetes on dried tomato halves was evidenced in duplicate by a two-step method involving olfactory (ie presence of off-flavours for yeasts) and visual (ie presence of mycelia for moulds) perception of the samples, followed by qualitative microbial analysis on a selective substrate.¹⁴ Microbial analyses were carried out by incubation under agitation of samples in enriched malt broth at 25 °C for 48 h. An aliquot of broth was then spread by a swab on a slide (Food Slide type 1, Pbi International, Milan, Italy) of Rose Bengal CAF agar. The slide was inserted into its tube, which was incubated at 30 °C for 48 h. Growth of moulds and yeasts was evidenced by visual perception of mycelia and colonies respectively.

This two-step method was able to show the following possible situations:

- positive sensory perception combined with a positive result from the microbial analysis showed that the storage conditions were able to grow eumycetes;
- negative sensory perception combined with a positive result from the microbial analysis showed that eumycetes were present on dried tomato halves,

but the storage conditions were not able to grow them;

- negative sensory perception combined with a negative result from the microbial analysis showed that eumycetes were not present on dried tomato halves since they had been inactivated during drying.

Response surface model

The following polynomial equation, describing the RSM of ΔE as a function of the variables studied, was set up:

$$\Delta E = \beta_0 + \beta_1 M + \beta_2 T + \beta_3 t + \beta_4 MT + \beta_5 Mt + \beta_6 Tt + \beta_7 M^2 + \beta_8 T^2 + \beta_9 t^2 \quad (3)$$

where M is the moisture content (%) of dried tomato halves, T is the storage temperature and t is the storage time.

Variables present in the linear terms represent the co-ordinates of the maximum value predicted; variables present in the quadratic terms represent the surface curvature; and the bifactorial cross-products represent the directions of axes of the geometric figure obtained by sectioning the surface area.

Effects of individual variables and their interactions were calculated by applying the Yates algorithm.⁹

Data processing

The multivariate statistical analysis to apply the saturated factorial design and to determine the RSM was carried out using the Unscrambler[®] version 7.01 (Camo AS, Trondheim, Norway) software package.

RESULTS AND DISCUSSION

Drying and storage tests

Fig 1 shows the kinetics of the different drying tests

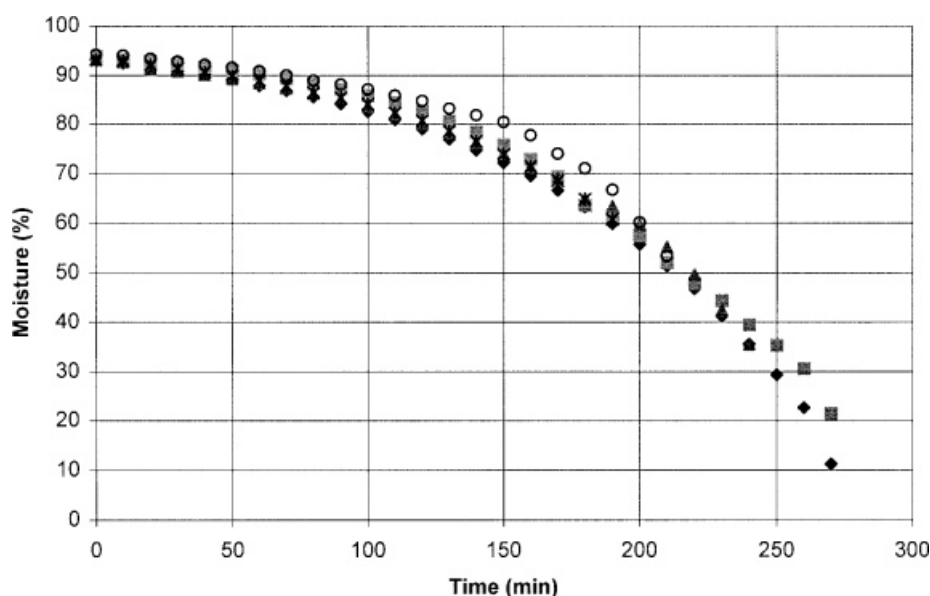


Figure 1. Moisture content (% on wet basis) of tomato halves versus drying time at 80 °C. Symbols refer to the final moisture contents: ◆, 10%; ■, 20%; ▲, 35%; ○, 50%; *, 60%.

Sample	Time	ΔE	<i>Eumycetes</i>
10% moisture ($a_w=0.45$)	$t=0$	—	—/—
	$t=19$ days at 17.5°C	4.18 ± 0.56	—/—
20% moisture ($a_w=0.69$)	$t=0$	—	—/+
	$t=8$ days at 10°C	5.55 ± 0.77	—/+
	$t=30$ days at 10°C	6.14 ± 0.63	—/+
	$t=8$ days at 25°C	11.36 ± 1.27	—/+
	$t=30$ days at 25°C	12.81 ± 1.52	—/+
35% moisture ($a_w=0.83$)	$t=0$	—	—/+
	$t=0.5$ day at 17.5°C	3.34 ± 1.90	—/+
	$t=19$ days at 17.5°C	4.04 ± 0.33	—/+
	$t=19$ days at 17.5°C	4.41 ± 0.23	—/+
	$t=37.5$ days at 17.5°C	10.06 ± 1.40	—/+
35% moisture ($a_w=0.83$)	$t=0$	—	—/+
	$t=19$ days at 5°C	1.78 ± 0.18	—/+
	$t=19$ days at 17.5°C	3.75 ± 0.46	—/+
	$t=19$ days at 17.5°C	3.89 ± 0.33	—/+
	$t=19$ days at 30°C	12.72 ± 1.57	—/+
50% moisture ($a_w=0.90$)	$t=0$	—	—/+
	$t=8$ days at 10°C	9.30 ± 0.53	—/+
	$t=30$ days at 10°C	7.83 ± 0.40	—/+
	$t=8$ days at 25°C	7.55 ± 0.39	—/+
	$t=30$ days at 25°C	19.58 ± 1.94	+/+
60% moisture ($a_w=0.93$)	$t=0$	—	—/+
	$t=19$ days at 17.5°C	11.53 ± 1.26	—/+

Table 3. Results from storage tests on samples at different moisture contents

carried out to reach the final moisture contents of dried tomato halves (ie 10, 20, 35, 50 and 60%) as defined by the saturated factorial design (Table 2).

Approximately 4 h and 30 min was required to remove water from tomato halves to 10% moisture and 3 h to remove water to 60% moisture. The different drying tests were carried out by a standard operating procedure, as evidenced by the similar kinetics.

Table 3 shows results from storage tests with different moisture contents of tomato halves. Moisture and water activity values for samples (a_w) determined by eqn (1) are also reported. It is evident that moisture contents include a wide range of water activity.

Results from microbial growth are marked with signs: the first sign refers to the olfactory and visual perception and the second sign to the microbial analysis. Apart from samples at 10 and 50% moisture, eumycetes were present in all samples, and their growth was inhibited by the storage conditions (ie —/+ signs in Table 3). At 10% moisture, eumycetes were not present (ie —/— signs in Table 3), suggesting that yeasts and moulds were inactivated during drying. At 50% moisture, growth of eumycetes was found on the sample stored at 25°C for 30 days (ie +/+ signs in Table 3), indicating that microbial growth was favoured by the storage conditions.

Optimisation by RSM

Table 4 shows the effects of individual variables and their interactions with respect to the RSM of ΔE (see eqn (3)).

It can be seen that storage temperature was the most important variable, whereas time and moisture content were less important. The minimum point for ΔE (ie 2.17) was predicted by the polynomial model as follows: storage of tomato halves with 27% moisture content at 10°C for 21 days. These conditions were tested experimentally by drying tomato halves to 30% final moisture, followed by storage at 10°C for 20 days. A measured value of $\Delta E = 3.25 \pm 0.26$ was obtained versus a calculated value of $\Delta E = 2.31$; the polynomial model was validated by the good reproducibility of the result.

The application of multiple regression analysis to ΔE data in Table 3 allowed us to determine response surfaces. Since these surfaces cannot be easily described, the relevant isoresponse curves are reported here (Figs 2–4).

Fig 2 shows the effect of temperature and time on

Table 4. Effects of individual variables and their interactions

Factor	Estimated effect
Average (β_0)	3.901
Moisture (β_1)	0.101
Temperature (β_2)	0.339
Time (β_3)	0.159
Moisture \times temperature (β_4)	−0.248
Moisture \times time (β_5)	0.855
Temperature \times time (β_6)	1.443
Moisture ² (β_7)	1.531
Temperature ² (β_8)	1.360
Time ² (β_9)	1.203

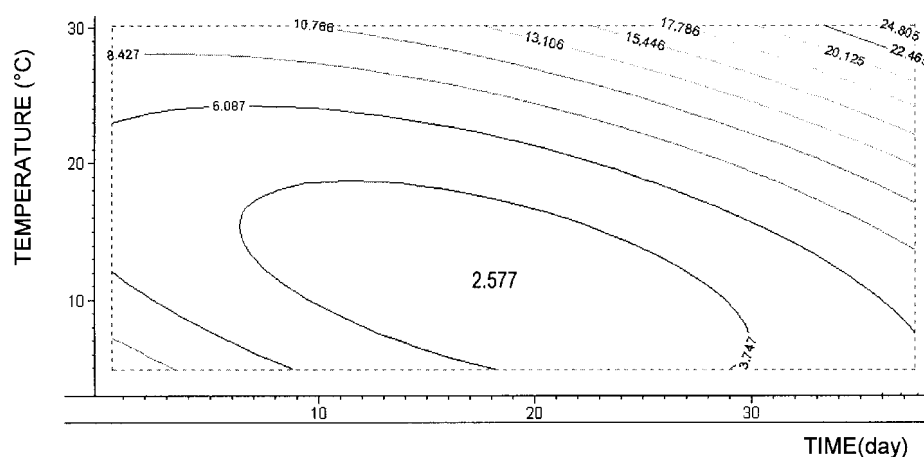


Figure 2. Isoresponse curves for ΔE as a function of temperature and time at 35% moisture content.

ΔE at the moisture content value of the centroid of the central composite design (ie 35%). A very small variation in ΔE (ie 2.58–6.09) occurred in dried tomato halves up to approximately 40 days of storage if the storage temperature was low (ie 5–10°C). At a storage temperature of 15–18°C the time required for minimum variation in ΔE decreased to approximately 30 days of storage.

Fig 3 shows the effect of moisture and time on ΔE at the storage temperature of the centroid of the central

composite design (ie 17.5°C). No considerable variation in ΔE (ie 3.28–6.09) occurred in dried tomato halves at 12–50% moisture content. The optimal range to minimise colour variation at the longest storage time (approximately 30 days) was 20–40% moisture content. Colour variation was also promoted at <12 and >50% moisture content.

Finally, Fig 4 shows the effect of moisture and temperature on ΔE at the storage time of the centroid of the central composite design (ie 19 days). No

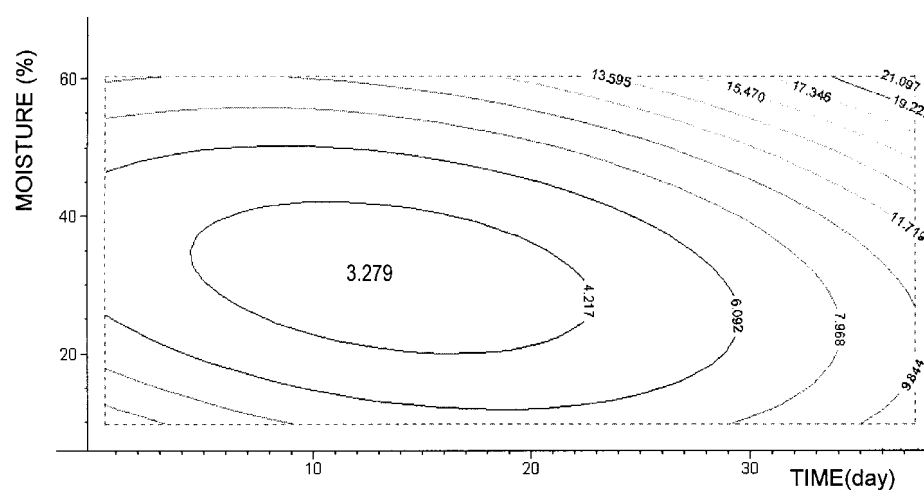


Figure 3. Isoresponse curves for ΔE as a function of moisture content and time at 17.5°C storage temperature.

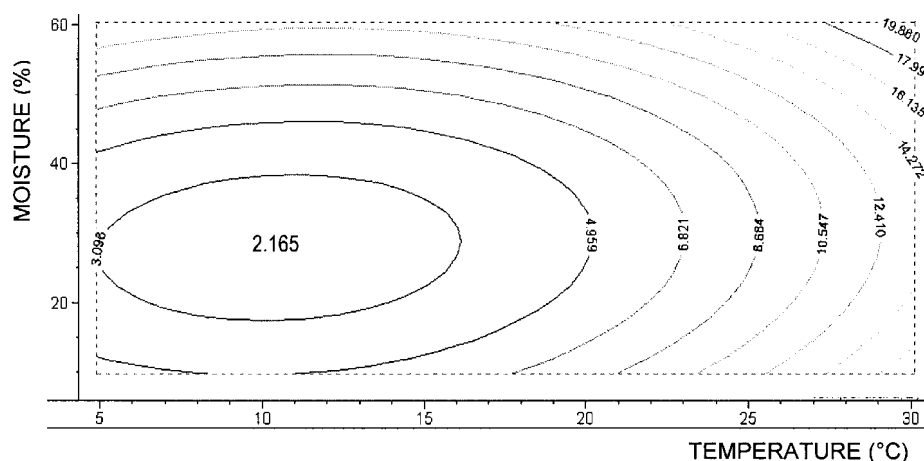


Figure 4. Isoresponse curves for ΔE as a function of moisture content and temperature at 19 days of storage time.

considerable variation in ΔE (ie 2.17–4.96) occurred in dried tomato halves at 10–45% moisture content and 5–20°C storage temperature. The range that allows one to minimise colour variation at storage temperatures close to room temperature was 20–40% moisture content.

Combining all the above results with the results on microbial growth described previously (Table 3), optimisation by RSM allowed us to define the following optimal storage operating conditions under vacuum in the dark to minimise colour variation of and, indirectly, oxidative damage to dried tomato halves:

- 20–40% moisture content, corresponding to $a_w = 0.69$ – 0.86 respectively, at $\leq 18^\circ\text{C}$ storage temperature, with a minimum point of colour variation at approximately 30% moisture content and 10°C .

CONCLUSIONS

Based on the results from this work, the following conclusions can be drawn.

Oxidative damage has to be minimised during storage of dried tomato to preserve sensory and nutritional characteristics of the product. Results show that temperature was the most important variable to minimise oxidative damage. In particular, at $\leq 18^\circ\text{C}$ storage temperature, oxidative damage to dried tomato halves can be minimised during storage under vacuum in the dark.

The moisture content of dried tomato halves also played an important role. The moisture content has to be reduced during drying to obtain adequately low water activity values in order to decrease the velocity of oxidative damage kinetics and to inhibit growth of spoilage micro-organisms. Interestingly, adequate shelf-life seems to be obtained at relatively high moisture content (ie 30–40%) and, consequently, at high water activity (0.80–0.86).

This conclusion should be combined with another observation: a low moisture content (ie $\leq 12\%$) of dried tomato halves seems to promote oxidative damage. This behaviour has been verified for lipid oxidation; when the moisture content falls below the monolayer moisture content of the product, an increase in oxidation can occur.¹⁵ The traditional production of dried tomato halves at low moisture content results in (i) significant oxidative heat damage due to both longer drying times and higher surface temperatures,¹ (ii) additional significant oxidative damage during storage and (iii) thermal death of eumycetes on the surface of tomato halves.

For tomato-drying optimisation, complete dehydration of the product should be avoided to minimise oxidative damage to the product. Hence the setting up of new tomato products at intermediate moisture may be interesting for the tomato industry. Pilot tests are being carried out to verify this hypothesis in terms of technological aspects, sensory acceptability and nutritional value of these new products.

ACKNOWLEDGEMENTS

We are grateful to R Nani of IVTPA (Milan, Italy) for his help in drying and storage tests and to C Peri of diSTAM for planning the work.

REFERENCES

- 1 Zanoni B, Peri C, Nani R and Lavelli V, Oxidative heat damage of tomato halves as affected by drying. *Food Res Int* **31**:395–401 (1999).
- 2 Cole ER and Kapur NS, The stability of lycopene. I. Degradation by oxygen. *J Sci Food Agric* **8**:360–365 (1957).
- 3 Cole ER and Kapur NS, The stability of lycopene. II. Oxidation during heating of tomato pulps. *J Sci Food Agric* **8**:366–368 (1957).
- 4 Khachik F, Goli MB, Beecher GR, Holden J, Lusby WR, Tenorio MD and Barrera MR, Effect of food preparation on qualitative and quantitative distribution of major carotenoids of tomatoes and several green vegetables. *J Agric Food Chem* **40**:390–398 (1992).
- 5 Sharma SK and Le Maguer M, Kinetics of lycopene degradation in tomato pulp solids under different processing and storage conditions. *Food Res Int* **29**:309–315 (1996).
- 6 Nicoli MC, Anese M, Parpinel MT, Franceschi S and Lerici CR, Loss and/or formation of antioxidants during food processing and storage. *Cancer Lett* **114**:71–74 (1997).
- 7 Anese M, Manzocco L, Nicoli MC and Lerici CR, Antioxidant properties of tomato juice as affected by heating. *J Sci Food Agric* **79**:750–754 (1999).
- 8 Baloch WA, Khan S and Baloch AK, Influence of chemical additives on the stability of dried tomato powder. *Int J Food Sci Technol* **32**:117–120 (1997).
- 9 Box GEP, Hunter WG and Hunter JS, *Statistics for Experimenters*. Wiley, New York (1978).
- 10 Alcaraz EC, Martin MA and Marin JP, Metodo manometrico para medida de humedades de equilibrio. *Grasas y Aceites* **28**:403–407 (1977).
- 11 Francis FJ and Clydesdale FH, *Food Colorimetry: Theory and Applications*. AVI Publishing, Westport, CT (1975).
- 12 D'Souza MC, Singha S and Ingle M, Lycopene concentration of tomato fruit can be estimated from chromaticity values. *HortSci* **27**:465–466 (1992).
- 13 Barreiro JA, Milano M and Sandoval AJ, Kinetics of colour change of double concentrated tomato paste during thermal treatment. *J Food Engng* **33**:359–371 (1997).
- 14 Stinson CG and Tiwari NP, Evaluation of quick bacterial count methods for assessment of food plant sanitation. *J Food Protect* **41**:269–271 (1978).
- 15 Labuza TP, Kinetics of lipid oxidation in foods. *CRC Crit Rev Food Technol* **2**:355–405 (1971).